AMENDMENTS TO THE CLAIMS:

This listing of the claims will replace all prior listings and versions of claims in the application.

CLAIMS:

- 1-31 (canceled)
- 32. (new) A method for analyzing microRNA, comprising:
 - a) contacting microRNA with a probe to form an RNA detection structure, wherein said probe comprises a first region that is complementary to said microRNA and a second region that is not complementary to said microRNA, wherein said second region can form a hairpin loop when said probe is hybridized to said microRNA; and
 - b) detecting said RNA detection structure.
- 33. (new) The method of claim 32, wherein said detecting comprises quantitating said microRNA.
- 34. (new) The method of claim 32, wherein said detecting comprises forming an invasive cleavage structure, cleaving said invasive cleavage structure, and detecting the cleavage of said invasive cleavage structure.
- 35. (new) The method of claim 32, wherein said detecting comprises use of a detection assay that employs sequence analysis.
- 36. (new) The method of claim 32, wherein said detecting comprises use of a detection assay that employs polymerase chain reaction.
- 37. (new) The method of claim 32, wherein said detecting comprises use of a detection assay that employs microarray hybridization.

- 38. (new) The method of claim 32, wherein said detecting comprises use of a detection assay that employs ligation.
- 39. (new) The method of claim 32, wherein said detecting comprises use of a labeled probe.
- 40. (new) The method of claim 39, wherein said labeled probe is fluorescently labeled.
- 41. (new) The method of claim 39, wherein said labeled probe is configured for FRET detection.
- 42. (new) The method of claim 41, wherein said labeled probe has a first conformation when not hybridized in a duplex and a second conformation when hybridized in a duplex.
- 43. (new) The method of claim 41, wherein said labeled probe exhibits increased fluorescence when hybridized in a duplex.
- 44. (new) The method of claim 32, wherein said detecting comprises use of a detection assay that employs polymerase chain reaction coupled with 5' nuclease cleavage of a labeled probe.
- 45. (new) The method of claim 44, wherein said labeled probe is fluorescently labeled.
- 46. (new) The method of claim 44, wherein said labeled probe is configured for FRET detection upon cleavage.

- 47. (new) The method of claim 32, wherein said detecting comprises exposing said RNA detection structure to a polymerase under conditions that permit primer extension.
- 48. (new) The method of claim 32, wherein said detecting comprises determining the presence of said microRNA in a sample.
- 49. (new) The method of claim 48, wherein said detecting comprises distinguishing said microRNA from another nucleic acid in said sample.
- 50. (new) The method of claim 49, wherein said sample comprises a cell lysate.
- 51. (new) The method of claim 32, wherein said microRNA is approximately 21-22 nucleotides in length.
- 52. (new) The method of claim 32, wherein a plurality of different microRNAs are detected.
- 53. (new) The method of claim 52, wherein said plurality of microRNAs comprise a first microRNA and a second microRNA that is said first microRNA having a polymorphism.
- 54. (new) The method of claim 32, wherein said microRNA is selected from the group consisting of Let-7, miR-1, miR-135, miR-15, miR-16, miR125b, miR-1d, and miR124a.
- 55. (new) The method of claim 32, wherein at least a portion of said RNA detection structure comprises a nucleotide analog.

56. (new) The method of claim 32, wherein at least a portion of said RNA detection structure comprises a peptide nucleic acid.